

IN THE CLAIMS

Please amend the claims as follows:

1-25 (Cancelled)

26. (Previously Presented) A method of preparing a metathesis catalyst, the method including the steps of:

mixing an aqueous solution of transition metal anions having a pH of 9 or higher with a carrier; and

removing water from the mixture by means of evaporation.

27. (Previously Presented) A method as claimed in Claim 26, wherein the carrier is silica and the transition metal is tungsten.

28. (Previously Presented) A method as claimed in Claim 27, wherein the aqueous solution contains tungsten in the form of ammonium metatungstatehydrate and/or ammonium tungstate.

29. (Currently Amended) A method as claimed in Claim ~~30~~ 28, wherein the aqueous solution contains tungsten in the form of ammonium metatungstatehydrate and wherein the concentration of the ammonium metatungstatehydrate and the mass of the silica are selected such that the $W O_3$ on the $S i O_2$ is from 4 to 10 wt%.

30. (Currently Amended) A method ~~of preparing a metathesis catalyst~~ as claimed in Claim 26, wherein excess water is removed by evaporation at about 80°C under reduced pressure to form a residue.

31. (Currently Amended) A method ~~of preparing a metathesis catalyst~~ as claimed in Claim 30, wherein further water is removed after removal of the excess water by drying the residue at about 110°C for about 12 hours, then by raising the temperature at a rate of about 1°C every minute up to about 250°C, maintained at about 250°C for about two hours and then by raising the temperature at a rate of about 3°C every minute up to about 550°C.

32. (Currently Amended) A method ~~of preparing a metathesis catalyst~~ as claimed in Claim 31, wherein the residue is then calcined.

33. (Currently Amended) A method ~~of preparing a metathesis catalyst~~ as claimed in Claim 32, wherein the residue is calcined at about 550°C for about 8 hours.

34. (Currently Amended) A method ~~of preparing a metathesis catalyst~~ as claimed in Claim 32, wherein the residue is calcined at a temperature and for duration such that the calcination step substantially removes NH_3 , ensures that the oxidation state of the tungsten is mostly 6+ and ensures that the tungsten oxide is bound to the carrier.

35. (Previously Presented) A catalyst for metathesis of an olefinic feed stream, which includes:
a transition metal oxide; and
a carrier, the transition metal oxide being deposited onto the carrier from an aqueous solution of transition metal oxide anions at a pH of 9 or more.

36. (Previously Presented) A catalyst as claimed in Claim 35, wherein the transition metal is tungsten and the carrier is silica.

37. (Previously Presented) A catalyst as claimed in Claim 35, wherein the catalyst is a heterogeneous catalyst.

38. (Previously Presented) A catalyst as claimed in Claim 36, wherein the catalyst is a heterogeneous catalyst.

39. (Previously Presented) A catalyst as claimed in Claim 37, wherein most of the tungsten oxide deposits are substantially amorphous.

40. (Previously Presented) A catalyst as claimed in Claim 38, wherein most of the tungsten oxide deposits are substantially amorphous.

41. (Previously Presented) A catalyst as claimed in Claim 36, wherein the catalyst is characterised in that at least a portion of some of the

tungsten oxide deposits are in the form crystallites of less than about 135 Å across on the surface of the carrier.

42. (Previously Presented) A catalyst as claimed in Claim 36, wherein the tungsten oxide is from about 4 to 10 wt% on SiO₂.

43. (Previously Presented) A catalyst as claimed in Claim 36, wherein the catalyst is characterised in that it remains catalytically active for at least 1000 hours at optimal operating conditions.

44. (Previously Presented) A catalyst as claimed in Claim 36, wherein the catalyst is characterised in that it provides a conversion rate of at least 30% for at least 50 hours at optimal operating conditions.

45. (Previously Presented) A metathesis process, which includes the step of:

contacting a C₅ and/or higher olefinic feed stream with a catalyst for metathesis at a temperature of between 350°C and 600°C, which catalyst includes:

a transition metal oxide; and

a carrier, the transition metal oxide being deposited onto the carrier from an aqueous solution of transition metal oxide anions at a pH of 9 or more.

46. (Previously Presented) A metathesis process as claimed in Claim 45, wherein the transition metal is tungsten and the carrier is silica.

47. (Previously Presented) A metathesis process as claimed in Claim 45, wherein the process includes a step of activating the catalyst at about 500 to 700°C for about 8 hours in an inert atmosphere.

48. (Previously Presented) A metathesis process as claimed in Claim 47, wherein the olefinic feed stream is selected such that the process yields C₁₀ to C₁₈ olefins.

49. (Previously Presented) A metathesis process as claimed in Claim 45, wherein the feed stream is contacted with the catalyst at a LHSV of between 5 and 25 h^{-1} at a temperature of between 350 and 550°C.

50. (Previously Presented) A metathesis process as claimed in Claim 45, wherein the feed stream is contacted with the catalyst at a pressure of 0.1 to 10 atm.